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CASE LA0108 NP

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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

IN RE APPLICATION OF

WASHBURN ET AL.

APPLICATION NO: 10/764,118

FILED: JANUARY 23, 2004

FOR: CYCLOALKYL CONTAINING ANILIDE LIGANDS FOR THE

THYROID RECEPTOR

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INFORMATION DISCLOSURE STATEMENT

Sir:

Applicants believe this paper is being filed before the mailing date of a first Office Action on the merits, and so under 37 C.F.R. §1.97(b)(3) no fees are required. If a fee is deemed to be required, the Commissioner is hereby authorized to charge such fee to Deposit Account No. 19-3880.

In accordance with 37 C.F.R. §1.56, applicants wish to call the Examiner's attention to the references cited on the attached form(s) PTO-1449.

Copies of these references are enclosed herewith.

The Examiner is requested to consider the foregoing information in relation to this application and indicate that each reference was considered by returning a copy of the initialed PTO 1449 form(s).

Bristol-Myers Squibb Company Patent Department P.O. Box 4000 Princeton, NJ 08543-4000 (609) 252-3816

Way 18,2004

Date:

Respectfully submitted,

Jonathan N. Provoost Attorney for Applicants Reg. No. 44,292

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

INFORMATION DISCLOSURE CITATION

E(Use several sheets if necessary)

ATTY. DOCKET NO. LA0108 NP APPLICATION NO. 10/764,118 APPLICANT WASHBURN ET AL. FILING DATE JANUARY 23, 2004

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U.S. PATENT APPLICATION PUBLICATIONS

EXAMINER INITIAL	_	U.S. APPLICATION DOCUMENT	DATE OF PUBLICATION	NAME	CLASS	SUBCLASS	FILING DATE
	AA	2002/0111315	8/15/02	Washburn et al.			

U.S. PATENT DOCUMENTS

EXAMINER INITIAL		DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	FILING DATE
	AB	3,239,345	3/8/66	Hodge et al.			
	AC	3,983,140	9/28/76	Endo et al.			
	AD	4,027,009	5/31/77	Grier et al.			
	AE	4,036,979	7/19/77	Asato			
	AF	4,231,938	11/4/80	Monaghan et al.			
	AG	4,346,227	8/24/82	Terahara et al.			
	АН	4,379,785	4/12/83	Weyer et al.			
	Al	4,411,890	10/25/83	Momany			
	AJ	4,448,784	5/15/84	Glamkowski et al.			

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	DOCUMENT NUMBER	DATE	OFFICE	CLASS	SUBCLASS	TRAN YES	SLATION NO
AK	EP 0 142 146	8/31/88	EP				
AL	EP 0 221 025	5/6/87	EP		,		
AM	EP 0 598 359	6/14/00	EP				
AN	EP 0 684 254	3/24/99	EP				
AO	EP 0 773 226	1/13/99	EP				

OTHER DOCUMENTS (Including Author, Title, Date, Pertinent pages, Etc.)

•	AP	Ashworth, D.M. et al., "2-Cyanopyrrolidides as Potent, Stable Inhibitors of Dipeptidyl Peptidase IV", Bioorganic & Medicinial Chemistry Letters, Vol. 6, No. 10, pp. 1163-1166 (1996)
	AQ	Ashworth, D.M. et al., "4-Cyanothiazolidides as Very Potent, Stable Inhibitors of Dipeptidyl Peptidase IV", Bioorganic & Medicinial Chemistry Letters, Vol. 6, No. 22, pp. 2745-2748 (1996)
	AR	Biller, S.A. et al., "Isoprenoid (Phosphinylmethyl)phosphonates as Inhibitors of Squalene Synthetase", Journal of Medicinal Chemistry, Vol. 31, No. 10, pp. 1869-1871 (1988)

EXAMINER

DATE CONSIDERED

^{*}EXAMINER: Initial of reference considered, whether or not citation is in conformance with MPEP 609: Draw a line through citation if not in conformance and not considered. Include a copy of this form with the next communication to applicant.

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EXAMINER INITIAL		DOCUMENT NUMBER	DATE	NAME	CL	ASS SUBC	LASS	FILING
	2AA	4,450,171	5/22/84	Hoffman et al.				DATE
	2AB	4,499,289	2/12/85	Baran et al.				
	2AC	4,572,912	2/25/86	Yoshioka et al.				
	2AD	4,613,610	9/23/86	Wareing				
<u> </u>	2AE	4,639,436	1/27/87	Junge et al.		.=		
	2AF	4,647,576	3/3/87	Hoefle et al.				
····	2AG	4,681,893	7/21/87	Roth				
	2AH	4,686,237	8/11/87	Anderson	· · · · ·	-		
	2AI	4,759,923	7/26/88	Buntin et al.				
	2AJ	4,871,721	10/3/89	Biller				
	2AK	4,904,769	2/27/90	Rauenbusch				
	2AL	4,924,024	5/8/90	Biller				
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			FOREIG	N PATENT DOCUMENTS				
		DOCUMENT NUMBER	DATE	OFFICE	CLASS	SUBCLASS	TRAN YES	ISLATIO NO
	2AM	EP 0 850 948	4/24/02	EP				
	2AN	EP 1 297 833	4/2/03	EP				
	2AO	2 596 393	10/2/87	FR				
	2AP	JP 8-27006	1/30/96	JP (with English abstract)				
	2AQ	JP 9-124684	5/13/97	JP (with English abstract)				
		OTHER DOC	UMENTS (Including Author, Title, Date, Pertino	ent nages F	itc)		•
	T			nase Inhibitors", Current Pharm			2 No.	1 nn 1
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		Bundgaard H Chapt	er 5: "Desig	n and Application of Prodrugs"	', A Textbo	ook of Drug D	esign pp. 11	and 3-191
	2AS	Development, Harwoo (1991)	od Academio	C Publishers, publ., Krogsgaard	Laroon, 1	. ct a, cas.,		
	2AS 2AT	Development, Harwoo (1991)		odrugs, Elsevier Science Publis				

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			U.S. F	PATENT DOCUMENTS			·	
EXAMINER INITIAL		DOCUMENT NUMBER	DATE	NAME	CL	ASS SUBC	LASS	FILING DATE
	3AA	5,006,530	4/9/91	Angerbauer et al.				DATE
*****	3AB	5,011,930	4/30/91	Fujikawa et al.				
-	3AC	5,177,080	1/5/93	Angerbauer et al.				<u> </u>
	3AD	5,260,440	11/9/93	Hirai et al.				
	3AE	5,273,995	12/28/93	Roth		· · · · · · · · · · · · · · · ·		
	3AF	5,346,701	9/13/94	Heiber et al.		-		
	3AG	5,354,772	10/11/94	Kathawala				
	3AH	5,385,929	1/31/95	Bjorge et al.				
	3AI	5,401,772	3/28/95	Yokoyama et al.				
	3AJ	5,488,064	1/30/96	Sher			Ì	
	3AK	5,491,134	⁻ 2/13/96	Sher et al.				
,	3AL	5,506,219	4/9/96	Robl				
			FOREIG	N PATENT DOCUMENTS				
		DOCUMENT NUMBER	DATE	OFFICE	CLASS	SUBCLASS	TRA	NSLATION NO
	3AM	JP 9-124685	5/13/97	JP (with English abstract)				
	3AN	JP 9-188625	7/22/97	JP (with English abstract)				
	3AO	JP 10-245391	9/14/98	JP				
	3AP	GB 2 205 837	12/21/88	UK				
	3AQ	WO 86/03488	6/19/86	PCT				
		OTHER DOC	UMENTS (Including Author, Title, Date, Pertine	ent pages, E	Etc.)		*****
	3AR	Squalene Biosynthesis	s", dissertati	uation of Ammonium Analogs on, Department of Medicinal C 8-51, Summary (June 1987)	of Carboc hemistry,	ationic Intern University of	nediate Utah,	es in pp. iv-v,
_	3AS	Chakrabartty, S.K., Ch	napter V: "A	lkaline Hypohalite Oxidations", ahanovsky, W.S., ed., pp. 343-	Oxidation 370 (1978	in Organic (Chemi	stry, Part
	3AT	Chan, D.M.T. et al., "N Tetrahedron Letters, \		D-Arylations with Phenylboroni 933-2936 (1998)	c Acids ar	nd Cupric Ace	etate",	
EXAMI	VER	<u> </u>		DATE CONSIDERED				

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		-	U.S. F	PATENT DOCUMENTS				
EXAMINER INITIAL		DOCUMENT NUMBER	DATE	NAME	CL	ASS SUBC	LASS	FILING DATE
	4AA	5,541,204	7/30/96	Sher et al.				
	4AB	5,594,016	1/14/97	Ueno et al.				
	4AC	5,595,872	1/21/97	Wetterau, II et al.				
	4AD	5,612,359	3/18/97	Murugesan				
	4AE	5,614,492	3/25/97	Habener				
	4AF	5,631,224	5/20/97	Efendic et al.				
· · ·	4AG	5,686,104	11/11/97	Mills et al.				
	4AH	5,691,322	11/25/97	Robl				
	4AI	5,712,279	1/27/98	Biller et al.				
	4AJ	5,712,396	1/27/98	Magnin et al.				-
	4AK	5,739,135	4/14/98	Biller et al.				
	4AL	5,753,675	5/19/98	Wattanasin				· · · · · · · · · · · · · · · · · · ·
			FOREIG	N PATENT DOCUMENTS				
, ,		DOCUMENT NUMBER	DATE	OFFICE	CLASS	SUBCLASS	TRAN	NSLATION NO
	4AM	WO 86/07054	12/4/86	PCT				
	4AN	WO 89/07110	8/10/89	PCT				
	4AO	WO 89/07111	8/10/89	PCT				
	4AP	WO 93/04081	3/4/93	PCT				
	4AQ	WO 96/38144	12/5/96	PCT	- 			
		OTHER DOC	UMENTS (Including Author, Title, Date, Pertine	nt pages, E	Etc.)	<u> </u>	
	4AR	Chiellini, G. et al., "A r Chemistry & Biology, '	nigh-affinity s Vol. 5, No. 6	subtype-selective agonist ligand , pp. 299-306 (1998)	for the th	yroid hormo	ne rec	eptor",
	4AS	Multistep Biosynthesis	 Demonstr 	Inreactive Analogs of Terpenoid ation That 'Presqualene Pyrophene", J. Am. Chem. Soc., Vol. 9	osphate'	Is an Essent	ial	
	4AT	Cornicelli, J.A. et al., "	15-Lipoxyge	nase and Its Inhibition: A Nove Design, Vol. 5, No. 1, pp. 11-20	l Therape			
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			U.S. I	PATENT DOCUMENTS				
EXAMINER INITIAL		DOCUMENT NUMBER	DATE	NAME	CL	ASS SUBC	LASS	FILING DATE
	5AA	5,760,246	6/2/98	Biller et al.				DATE
	5AB	5,770,615	6/23/98	Cheng et al.				
	5AC	5,776,983	7/7/98	Washburn et al.				
	5AD	5,827,875	10/27/98	Dickson, Jr. et al.				-
	5AE	5,885,983	3/23/99	Biller et al.				
	5AF	5,962,440	10/5/99	Sulsky				
	5AG	6,043,265	3/28/00	Murugesan et al.				
	5AH	6,184,231	2/6/01	Hewawasam et al.				
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		DOCUMENT NUMBER	DATE	OFFICE	CLASS	SUBCLASS	TRAN YES	NSLATION - NO
	5AM	WO 97/12613	4/10/97	PCT				
	5AN	WO 97/12615	4/10/97	PCT				
	5AO	WO 97/21993	6/19/97	PCT				
	5AP	WO 99/00353	1/7/99	PCT		·		
	5AQ	WO 99/38501	8/5/99	PCT		•		
		OTHER DOC	UMENTS (Including Author, Title, Date, Pertine	nt pages, E	tc.)		
	5AR	Couladouros, E.A. et a eastern part of bastad	al., "A gener ins 4-16", Te	al synthetic route towards basta etrahedron Letters, Vol. 40, pp.	dins. Pa 7023-702	rt 1: Synthe: 6 (1999)	sis of t	he
	5AS			Thyroxine and Related Compouted to Thyroxine", J. Chem. Soc				ration of
	5AT	Driver, M.S. et al., "A Amines from Aryl Hali 118, No. 30, pp. 7217	des and Prir	<u> </u>	Amination PF)PdCl ₂ "	n: Mixed Se , J. Am. Che	condar m. So	c., Vol.
EXAMI	NFR			DATE CONSIDERED				

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	DOCUMENT NUMBER	DATE	OFFICE	CLASS	SUBCLAS	TRAN YES	SLATION NO
6AA	WO 99/46272	9/16/99	PCT				
6AB	WO 99/61431	12/2/99	PCT				
6AC	WO 99/67278	12/29/99	PCT		·		
6AD	WO 99/67279	12/29/99	PCT				
6AE	WO 00/01389	1/13/00	PCT				
6AF	WO 01/60784	8/23/01	PCT				
6AG	WO 01/70687	9/27/01	PCT				
6AH	WO 01/72692	10/4/01	PCT				
6AI	WO 01/85670	11/15/01	PCT				
6AJ	WO 01/90053	11/29/01	PCT				
6AK	WO 01/94293	12/13/01	PCT				
6AL	WO 02/051805	7/4/02	PCT				
6AM	WO 02/062780	8/15/02	PCT				
6AN	WO 02/090344	11/14/02	PCT				
6AO	WO 02/094319	11/28/02	PCT				
6AP	WO 2004/018421	3/4/04	PCT				
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EXAMINER	DATE CONSIDERED
7AN	Horner, L. et al., "Die Synthese brücken-analoger Thyroninverbindungen", Chemische Berichte, Vol. 85, pp. 520-530 (1952)
7AM	Hongu, M. et al., "Na*-Glucose Cotransporter Inhibitors as Antidiabetic Agents. III. Synthesis and Pharmacological Properties of 4'-Dehydroxyphlorizin Derivatives Modified at the OH Groups of the Glucose Moiety", Chem. Pharm. Bull., Vol. 46, No. 10, pp. 1545-1555 (1998)
7AL	Hongu, M. et al., "Na ⁺ -Glucose Cotransporter Inhibitors as Antidiabetic Agents. II. Synthesis and Structure-Activity Relationships of 4'-Dehydroxyphlorizin Derivatives", Chem. Pharm. Bull., Vol. 46, No. 1, pp. 22-33 (1998)
7AK	Hickey, D.M.B. et al., "Synthesis of Thyroid Hormone Analogues. Part 3. Iodonium Salt Approaches to SK&F L-94901", J. Chem. Soc. Perkin Trans. I, pp. 3103-3111 (1988)
7AJ	Hickey, D.M.B. et al., "Synthesis of Thyroid Hormone Analogues. Part 2. Oxidative Coupling Approach to SK&F L-94901", J. Chem. Soc. Perkin Trans. I, pp. 3097-3102 (1988)
7AI	Harrington, C.R., "Synthesis of a Sulphur-containing Analogue of Thyroxine", Biochem. J., Vol. 43, pp. 434-437 (1948)
7AH	Hara, S., "Ileal Na ⁺ /bile acid cotransporter inhibitors", Drugs of the Future, Vol. 24, No. 4, pp. 425-430 (1999)
7AG	Hamann, L.G. et al., "Discovery of a Potent, Orally Active, Nonsteroidal Androgen Receptor Agonist: 4-Ethyl-1,2,3,4-tetrahydro-6-(trifluoromethyl)-8-pyridono[5,6-g]-quinoline (LG121071)", J. Med. Chem., Vol. 42, No. 2, pp. 210-212 (1999)
7AF	Guo, ZW. et al., "Enzymatic Oxidative Phenolic Coupling", J. Org. Chem., Vol. 62, No. 20, pp. 6700-6701 (1997)
7AE	Greene, T.W. et al., Protective Groups in Organic Synthesis, Third Edition, John Wiley & Sons, Inc., publ. (1999) (table of contents)
7AD	Ghiselli, G., "The Pharmacological Profile of FCE 27677: A Novel ACAT Inhibitor with Potent Hypolipidemic Activity Mediated by Selective Suppression of the Hepatic Secretion of ApoB-100-Containing Lipoprotein", Cardiovascular Drug Reviews, Vol. 16, No. 1, pp. 16-30 (1998)
7AC	Frost, C.G. et al., "Recent developments in aromatic heteroatom coupling reactions", J. Chem. Soc., Perkin Trans. 1, pp. 2615-2623 (1998)
7AB	Evans, D.A. et al., "Synthesis of Diaryl Ethers through the Copper-Promoted Arylation of Phenols with Arylboronic Acids. An Expedient Synthesis of Thyroxine", Tetrahedron Letters, Vol. 39, pp. 2937-2940 (1998)
7AA	Edwards, J.P. et al., "Nonsteroidal Androgen Receptor Agonists Based on 4-(trifluoromethyl)-2 <i>H</i> -pyrano[3,2- <i>g</i>]quinolin-2-one", Bioorganic & Medicinal Chemistry Letters, Vol. 9, pp. 1003-1008 (1999)
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*EXAMINER: Initial of reference considered, whether or not citation is in conformance with MPEP 609: Draw a line through citation if not in conformance and not considered. Include a copy of this form with the next communication to applicant.